FDA 510(k) Premarket Notification

K060247 LIFE SPINE CEMENT RESTRICTOR

CLEARED

May 11, 2006
100 days to decision

K060247 · Product code: **JDK** · Orthopedic
Source: https://www.510kdatabase.net/k060247/

SUBMISSION DETAILS

Decision	Substantially Equivalent - U
Submission type	Traditional
Device classification	Prosthesis, Hip, Cement Restrictor (JDK)
Date received	Jan 31, 2006
Decision date	May 11, 2006
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Life Spine**
Location	Hoffman Estates, IL, US
Contact	ERIN MALLOY
510(k) history	36 submissions · 34 cleared · 2006-2018